PCT/KR2003/001494

PATENT COOPERATION TREATY PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Artcle 36 and Rule 70)

Applicant's or agent's file reference 2003OPA2765 // 44 aply KR	FOR FURTHER ACTION SeeNotificationofTransmittalofInternation Examination Report (Form PCT/IPEA/410			
International application No. PCT/KR2003/001494	International filing date(day/m 25 JULY 2003 (25.07.20		Priority date (day/month/ye	ar)
International Patent Classification (IPC)	or national classification and IF	c		
IPC7 C07C 29/00				
Applicant	······································	ı		
POSTECH Foundation et al				
This international preliminary examd is transmitted to the applicant. This REPORT consists of a total. This report is also accomp.	t according to Article 36. of sheets, included by ANNEXES, i.e., sheets	ding this cover she	et. , claims and/or drawings wl	hich have been
amended and are the basis	for this report and/or sheets core Administrative Instructions un	itaining rectificatio	ns made before this Autho	rity (see Rule
These annexes consist of a total	of sheets.			
3. This report contains indications relating to the following items: 1				
Date of submission of the demand	Date	of completion of t	this report	
25 FEBRUARY 2005	(25.02.2005)	23 SEPTEMB	BER 2005 (23.09.2005)	
Name and mailing address of the IPEA/ Korean Intellectual Property 920 Dunsan-dong, Seo-gu, Republic of Korea	y Office	norized officer MOON, Sun Heu	up	4/10
Facsimile No. 82-42-472-7140	Tele	phone No. 82-42	-481-5543	And pass

Form PCT/IPEA/409 (cover sheet) (July 1998)

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/KR2003/001494

1.	Basis	of the report	
1.	With	regard to the elements of the international application:*	
	\boxtimes	the international application as originally filed	
	\boxtimes	the description:	, as originally filed
		pages 1- 22 pages NONE	, filed with the demand
		pages NONE , filed with the letter of	
	\boxtimes	the claims:	
	نے	pages 23 - 31 pages NONE , as amended (together with any	, as originally filed statment) under Article 19
		pages NONE	en n 1.1 .1 .1 .1
		pages, filed with the letter of	, · · · · · · · · · · · · · · · · · · ·
		the drawings:	an aminimally filed
		pages	, as originally filed , filed with the demand
İ		pages filed with the letter of	· · · · · · · · · · · · · · · · · · ·
		the sequence listing part of the description:	, as originally filed
		pagespages	, filed with the demand
		pages, filed with the letter of	
2.	the i	regard to the language, all the elements marked above were available or furnished to this Authoritemational application was filed, unless otherwise indicated under this item. the elements were available or furnished to this Authority in the following language	shwhich is .1(b)).
3.	Wit prel	h regard to any nucleotide and/or amino acid sequence disclosed in the international appli iminary examination was carried out on the basis of the sequence listing: contained inthe international application in written form. filed together with the international application in computer readable form. furnished subsequently to this Authority in written form.	ication, the international
	끔	furnished subsequently to this Authority in computer readable form	
		The statement that the subsequently furnished written sequence listing does not go be international applicationas as filed has been furinshed. The statement that the information recorded in computer readable form is identical to the volume been furnished.	
4.		The amendments have resulted in the cancellation of: the description, pages the claims, Nos.	
5.		This report has been established as if (some of) the amendments had not been made, since go beyond the disclosure as filed, as indicated in the Supplemental Box(Rule 70.2(c)).**	they have been considered to
•	Repla in this and 7	cement sheets which have been furnished to the receiving Office in response to an invitation un opinion as "originally filed." and are not annexed to this report since they do not contain 0.17).	der Article 14 are referred to amendments (Rules 70,16
**	Any r	eplacement sheet containing such amendments must be referred to under item I and annexed to	o this report.

INTERNATIONAL PRELIMINARY EXAMINATION

International application No.
PCT/KR2003/001494

v. Reasoned statement under Article 35(2) with regard	to novelty, inventive step or industrial applicability;
citations and explanations supporting such statemen	<u> </u>
1	

Statement		1 - 21_	YES
Novelty (N)	Claims - <u> · · · · · · · · · · · · · · · ·</u>		
	Claims	NONE	N0
Inventive step (IS)	Claims	1 - 21	YES
• ` `	Claims	NONE	NO
Industrial applicability (IA)	Claims	1 - 21	YES
	Claims	NONE	_NO

- 2. Citations and explanations (Rule 70.7)
 - 1. Reference is made to the following documents:

D1: KR 2001-0040121

D2: Organometallics(1999, v.18, PP.3981-3990)

2. Novelty and Inventive Step

The present invention relates to a method for preparing (S)-chiral alcohol with high yield and high optical purity by mixing achiral substrates such as racemic alcohol or ketone with metal catalyst and protein hydrolase to perform a dynamic kinetic resolution reaction.

Document D1, which is considered to represent the most relevant state of the art, discloses a process for preparing a chiral ester by reacting a)racemic alcohol, b)a ruthenium complex catalyst, c)a lipase to acylate selectively one of enantiomers of said racemice alcohol, and d)an acyl donor group to supply acyl group to said lipase.

Docment D2 discloses the racemization of α -hydroxy ester using Pseudomonas cepacia lipase, ruthenium catalyst, and 4-chlorophenyl acetate as acyl donor in cyclohexane.

Although D1-D2 relate to methods for preparing optically active alcohol using enzyme catalyst, metal catalyst, and acyl donor like the present invention, they are different from the present invention in that since lipase is used as enzyme catalyst in the prior art documents, only R-entiaomer(that is, R-chiral alcohol) can be synthesized, whereas the present invention can provide a method of synthesizing (S)-chiral alcohol enantioselectively with high purity and high yield.

Moreover, a protein hydrolysis enzyme in the present invention, which plays a useful role in stimulating the enantioselective acylation of a racemic compound, is used as opposite stereoselectivity to lipase in D1-D2. Consequently, the present invention provides a novel process for preparing the (S)-chiral alcohol which is not easily exchangeable by the those who are skilled in the art and suggests a synthesis method of (S)-chiral alcohols with high optical purity and high yield.

Therfore, the subject matter of claims 1-21 of the present invention is considered to be novel and to involve an inventive step in the sense of PCT Article 33(2) and (3).

3. Industrial applicability

The subject matter of the claim 1-21 is considered to be industrially applicable under PCT Article 33(4).